

13. Motew SJ, Cherr GS, Craven TE, Travis JA, Wong JM, Reavis SW, et al. Renal duplex sonography: main renal artery versus hilar analysis. *J Vasc Surg* 2000;32:462-71.
14. Olin JW, Piedmonte MR, Young JR, DeAnna S, Grubb M, Childs MB. The utility of duplex ultrasound scanning of the renal arteries for diagnosing significant renal artery stenosis. *Ann Intern Med* 1995;122:833-8.
15. Avasthi PS, Voyles WF, Greene ER. Noninvasive diagnosis of renal artery stenosis by echo-Doppler velocimetry. *Kidney Int* 1984;25:824-9.
16. Hoffmann U, Edwards JM, Carter S, Goldman ML, Harley JD, Zaccardi MJ, et al. Role of duplex scanning for the detection of atherosclerotic renal artery disease. *Kidney Int* 1991;39:1232-9.
17. Miralles M, Cairós M, Cotillas J, Giménez A, Santiso A. Value of Doppler parameters in the diagnosis of renal artery stenosis. *J Vasc Surg* 1996;23:428-35.
18. Oliva VL, Soulez G, Lesage D, Nicolet V, Roy MC, Courteau M, et al. Detection of renal artery stenosis with Doppler sonography before and after administration of captopril: value of early systolic rise. *AJR Am J Roentgenol* 1998;170:169-75.
19. Strotzer M, Fellner CM, Geissler A, Gmeinwieser J, Kohler SM, Krämer BK, et al. Noninvasive assessment of renal artery stenosis. A comparison of MR angiography, color Doppler sonography, and intraarterial angiography. *Acta Radiol* 1995;36:243-7.
20. Kaplan-Pavlovic S, Nadja C. Captopril renography and duplex Doppler sonography in the diagnosis of renovascular hypertension. *Nephrol Dial Transplant* 1998;13:313-7.
21. Riehl J, Fritz A, Sieberth HG. The use of duplex sonography in the diagnosis of renal artery stenosis. *Eur J Med Res* 1997;2:14-22.
22. Berland LL, Koslin DB, Routh WD, Keller FS. Renal artery stenosis: prospective evaluation of diagnosis with color duplex US compared with angiography. *Work in progress. Radiology* 1990;174:421-3.
23. Edwards JM, Zaccardi MJ, Strandness DE Jr. A preliminary study of the role of duplex scanning in defining the adequacy of treatment of patients with renal artery fibromuscular dysplasia. *J Vasc Surg* 1992;15:604-9; discussion: 609-11.
24. Fleming SH, Davis RP, Craven TE, Deonanan JK, Godshall CJ, Hansen KJ. Accuracy of duplex sonography scans after renal artery stenting. *J Vasc Surg* 2010;52:953-7; discussion: 958.
25. Hua HT, Hood DB, Jensen CC, Hanks SE, Weaver FA. The use of colorflow duplex scanning to detect significant renal artery stenosis. *Ann Vasc Surg* 2000;14:118-24.
26. Olin JW. Role of duplex ultrasonography in screening for significant renal artery disease. *Urol Clin North Am* 1994;21:215-26.
27. Zierler RE, Bergelin RO, Isaacson JA, Strandness DE, Jr. Natural history of atherosclerotic renal artery stenosis: a prospective study with duplex ultrasonography. *J Vasc Surg* 1994;19:250-7; discussion: 257-8.
28. Eklöf H, Ahlström H, Magnusson A, Andersson LG, Andrén B, Hägg A, et al. A prospective comparison of duplex ultrasonography, captopril renography, MRA, and CTA in assessing renal artery stenosis. *Acta Radiol* 2006;47:764-74.
29. Solar M, Zizka J, Krajina A, Michl A, Raupach J, Kizo L, et al. Comparison of duplex ultrasonography and magnetic resonance imaging in the detection of significant renal artery stenosis. *Acta Med (Hradec Kralovce)* 2011;54:9-12.
30. Clemente A, Macchi V, Porzionato A, Stecco C, De Caro R, Morra A. CTA and 2D-3D post-processing: radiological signs of fibromuscular dysplasia of renal artery. *Surg Radiol Anat* 2009;31:25-9.
31. Martin RL, Nanra RS, Włodarczyk J, Desilva A, Bray A. Renal hilar Doppler analysis in the detection of renal artery stenosis. *J Vasc Technol* 1991;15:173-80.
32. Stavros AT, Parker SH, Yakes WF, Chantelois AE, Burke BJ, Meyers PR, et al. Segmental stenosis of the renal artery: pattern recognition of tardus and parvus abnormalities with duplex sonography. *Radiology* 1992;184:487-92.
33. Isaacson JA, Zierler RE, Spittell PC, Strandness DE. Noninvasive screening for renal artery stenosis: comparison of renal artery and renal hilar duplex scanning. *J Vasc Technol* 1995;19:105-10.
34. Greene ER, Venters MD, Avasthi PS, Conn RL, Jahnke RW. Noninvasive characterization of renal artery blood flow. *Kidney Int* 1981;20:523-9.
35. Bardelli M, Jensen G, Volkmann R, Aurell M. Non-invasive ultrasound assessment of renal artery stenosis by means of gosling pulsatility index. *J Hypertens* 1992;10:958-9.
36. Schwerk WB, Restrepo IK, Stellwaag M, Klose KJ, Schade-Brittinger C. Renal artery stenosis: grading with image-directed Doppler US evaluation of renal resistive index. *Radiology* 1994;190:785-90.
37. Zoller WG, Hermans H, Bogner JR, Hahn D, Middeke M. Duplex sonography in the diagnosis of renovascular hypertension. *Klin Wochenschr* 1990;68:830-4.
38. Kawarada O, Yokoi Y, Takemoto K, Morioka N, Nakata S, Shiotani S. The performance of renal duplex ultrasonography for the detection of hemodynamically significant renal artery stenosis. *Catheter Cardiovasc Interventions* 2006;68:311-8.
39. de Haan MW, Kroon AA, Flobbe K, Kessels AG, Tordoir JH, van Engelsehoven JM, et al. Renovascular disease in patients with hypertension: detection with duplex ultrasound. *J Hum Hypertens* 2002;16:501-7.
40. Strandness DE Jr. Duplex imaging for the detection of renal artery stenosis. *Am J Kidney Dis* 1994;24:674-8.
41. Cohn EJ Jr, Benjamin ME, Sandager GP, Lilly MP, Killewich LA, Flinn WR. Can intrarenal duplex waveform analysis predict successful renal artery revascularization? *J Vasc Surg* 1998;28:471-80; discussion: 480-1.

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DISCUSSION

Dr Kimberley Hansen (*Winston Salem, NC*). Dr Mitchell, Dr Endean, members, and guests: Thank you for the opportunity to open this discussion, and thank you Ali for your paper well in advance and your presentation this morning. For me, the primary message from Dr AbuRahma and his group is that optimal criteria for renal duplex sonography are best obtained when each laboratory validates its results through comparative analysis. We agree wholeheartedly with this message, but the comparison may not be so easy to make as it once was. Some years ago, our first comparative analysis between renal duplex and angiography utilized cut film, not temporal digital subtraction. Digital subtraction with postprocessing—peak opacification and pixel shift functions—can affect images profoundly. With an analog-to-digital imaging system, our technologists can create just about any image you might prefer, bringing me to these questions:

1. Who created the final subtracted angiogram for comparison?
2. Do your angiograms reflect analog-to-digital or digital-to-digital technologies?
3. In the absence of poststenotic dilatation or collaterals, how confident are you that 50, 60 and 70% lesions can be delineated one from the other?

More than any particular velocity cut point, the significance of a renal artery occlusive lesion depends on the clinical setting in which it occurs. In this regard, our group has considered severe hypertension as the clinical hallmark of physiologically significant renovascular disease, leading to these questions:

4. The majority of patients were considered to be hypertensive. How many patients had severe hypertension of the variety most consistent with a renovascular etiology?

5. Did the severity of hypertension vary with renal artery peak systolic velocity?

One last point: Velocity criteria for renal duplex will likely vary from laboratory to laboratory, but the added value of renal aortic ratio is uncertain. When Gene Strandness and Ted Kohler introduced renal aortic ratio in the early and mid-1980s, the supporting rationale was that velocities in the renal artery were dependent on inflow aortic velocity. In reality, renal blood flow is determined by autoregulation provided by the kidney. In large groups of volunteers, as well as patients, there is no association between peak aortic and peak renal artery velocities. For us, renal aortic ratio has proved to be a spurious correlation. The correlation with disease has rested entirely with renal artery peak systolic velocity. I enjoyed this paper and thank the association for the privilege of the floor.

Dr Ali F. AbuRahma. Thank you, Kim, I appreciate your constructive comments, and I am honored that you were the discussant for this paper. All of us are aware of your contributions on this subject, and they are appreciated. In regards to questions 1 and 2, the technical staff completed the digital subtraction angiograms and they were digital-to-digital, which is compatible with

most modern technology. In regards to question 3, as to how confident we were in measuring 50%, 60%, or 70%, we were certainly confident, particularly since our measurements were based on either $\geq 50\%$ or $>60\%$, and we did not look to $>70\%$, since it has no practical implication, at least at this stage. Two observers made the determination and if there was a difference of $\geq 10\%$, a third observer was selected and a consensus was reached. In regards to question 4, as noted in our presentation, over 90% of the patients were hypertensive and a majority were labeled as having severe hypertension (ie, a diastolic blood pressure of over 100 or a systolic blood pressure above 180); however, we did not classify patients according to the degree of hypertension.

In regards to the severity of hypertension and its relation to peak systolic velocity, as indicated earlier, we have not classified this accordingly; however, we are aware of the fact that velocities can vary in patients with cardiovascular disease. Therefore, we analyzed not only peak systolic velocities, but also the renal-aortic systolic ratio, and we found that a ratio of ≥ 3.7 was somewhat equivalent in its value to a peak systolic velocity of 285. Again, thank you for discussing this paper.